

## Metabolic adaptation and hormonal regulation in young rabbit does during long-term caloric restriction and subsequent compensatory growth

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### Abstract

*An experiment was performed to assess the metabolic adaptation and hormonal regulation in young female rabbits during long-term food restriction and subsequent compensatory growth during rearing. Feeding level was either ad libitum (AL, no. = 52) or restricted (R, no. = 52). From 6 to 12 weeks of age, food intake of R was kept at a constant level. This resulted in an increase in relative restriction as compared with AL to 0.54 of AL intake at 12 weeks of age (restriction period). Thereafter food intake gradually increased to 0.95 of AL at 17 weeks of age (recovery period). During the last 5 days before insemination at 17.5 weeks of age, all animals were fed to appetite. Blood samples were taken weekly from 6 to 17 weeks of age from 11 animals in each group. Growth rate of R was reduced during the restricted period (29 (s.d. 2) v. 44 (s.d. 5) g/day for R and AL, respectively;  $P < 0.05$ ), but was higher in the recovery period (30 (s.d. 3) v. 27 (s.d. 4) g/day, respectively;  $P < 0.05$ ). At first insemination, AL rabbits were heavier than R (4202 (s.d. 388) v. 3798 (s.d. 220) g, respectively;  $P < 0.001$ ). During the restricted period, plasma glucose was constantly lower ( $P < 0.05$ ) in R. Insulin levels paralleled those of glucose, being lower ( $P < 0.05$ ) in R than in AL. Restriction reduced ( $P < 0.05$ ) circulating corticosterone and tri-iodothyronine (T3) levels in R. Leptin, non-esterified fatty acids, and plasma urea nitrogen levels were similar for AL and R during food restriction, whereas triglycerides were similar until 10 weeks of age, after which the levels were lower in R. During the recovery period, the food intake of the R but not AL rabbits increased. Insulin was the only hormone in R rabbits that had returned to levels found in AL rabbits by the 2nd week of the recovery period. Glucose, T3, and corticosterone levels returned to levels found in AL rabbits between 3 to 4 weeks after refeeding. Non-esterified fatty acids, triglycerides, and leptin were higher ( $P < 0.05$ ) in AL rabbits from 13 weeks of age onwards. The pattern of changes in the endocrine status during food restriction and compensatory growth in rabbits do conform with those from other species, although some specific changes may vary depending on the severity of food restriction and its duration.*

**Keywords:** energy consumption, hormones, leptin, rabbits.

## Introduction

In commercial rabbit production, young rabbit does are given food *ad libitum* until first mating, which commonly occurs between 14 and 16 weeks of age when 0.75 to 0.80 of final body weight is reached (Lebas *et al.*, 1986). However, with this rearing strategy does are not mature enough for reproduction. In fact, both lower reproductive performance and high replacement rate do occur, probably due to their energy deficit during first lactation (Xiccato, 1996). Voluntary food intake is considered the main limiting factor that cannot be adjusted sufficiently by management measures (Cervera *et al.*, 1993; Parigi-Bini *et al.*, 1996). Delaying first mating to older age results in more mature, but fatter does, in which kit mortality at kindling is increased (Partridge *et al.*, 1986).

Recently, a different way to cope with the energy losses has been suggested by Rommers *et al.* (1999) in which the feeding strategy during rearing is adapted to delay sexual maturity and to stimulate body reserves and food intake capacity of young rabbit does (Rommers *et al.*, 2001a, 2001b and 2004). Although the effects of long-term food restriction and refeeding on the regulatory mechanisms of metabolism during reduced and compensatory growth have been investigated in several animal species (Yambayamba *et al.*, 1996; Rhind *et al.*, 2000), the adjustments of metabolic hormones during long-term food restriction and refeeding have not been investigated in rabbits. Therefore, an experiment was performed to assess the metabolic adaptation and hormonal regulation in young rabbit does during long-term food restriction and subsequent compensatory growth by examining the time course of several key hormones, such as leptin, insulin, triiodothyronine (T3), corticosterone, and intermediate metabolites from early post weaning to first insemination at 17.5 weeks of age.

## Material and methods

The Use and Care Committee of the Animal Sciences Group of Wageningen University and Research Center approved all protocols.

### Animals and husbandry

Young New Zealand White rabbit does (no. = 104) bred at Applied Research, were used in the experiment. Before weaning, kits were raised in litters of nine kits. At weaning (30 days of age), from each litter, pairs of female kits with live weight deviating by less than 15% of the average litter weight, were randomly taken. Sisters were put into adjacent individual cages (50 × 60 × 30 cm) of galvanized wire net, equipped with an automatic drinker and a manual feeder. After weaning, animals

were offered a standard commercial diet, containing 10.3 MJ/kg metabolizable energy (ME) and 170 g/kg crude protein (ABCETA, Lochem, The Netherlands). From 6 weeks of age onwards, animals were given food according to treatment until the end of the experiment. The experiment ended at first insemination (end of rearing), at 17.5 weeks of age.

Animals were housed in identical deep-pit compartments with controlled lighting. Before weaning, animals were kept at a 16-h photoperiod and a minimum ambient temperature of 16°C. During rearing, animals were submitted to a photoperiod of 12 h. The first 2 weeks after weaning, a minimum ambient temperature of 18°C was maintained. From 2 weeks after weaning onwards, the minimum ambient temperature was set at 16°C.

### Treatments

Treatments started at 6 weeks of age when sisters were randomly assigned to one of the two treatments. Food was provided *ad libitum* (AL, no. = 52) or restricted (R, no. = 52). The R rabbits were offered daily a restricted amount of food, aiming to follow a fixed body weight curve (see Figure 1) as described in a previous experiment (Rommers *et al.*, 2001a). From 6 to 12 weeks of age, the food intake was kept between 92 and 110 g/day (see Figure 2). This resulted in an increase in relative restriction as compared with AL to 0.54 of AL intake at 12 weeks of age (restriction period). From 12 weeks of age onwards, food intake was increased from 115 (s.d. 4) g/day to 192 (s.d. 14) g/day (0.95 of AL) at 17 weeks of age (recovery period, see Figure 2). In the AL rabbits voluntary food intake increased until 13 weeks of age. From this age onwards, food intake remained at approximately 200 g/day.

During the last 5 days before insemination (at 17.5 weeks of age), all animals were given food to appetite. R rabbits consumed 235 (s.d. 39) g and

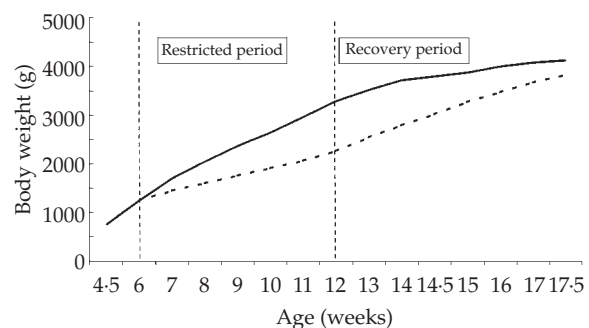
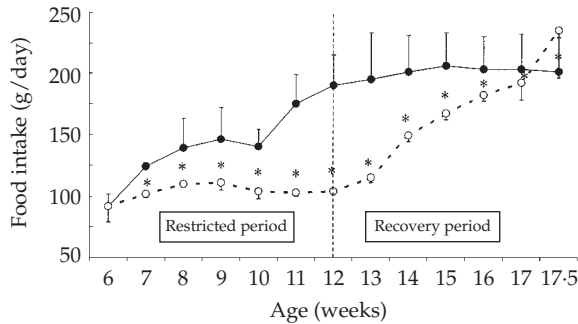


Figure 1 Fixed body weight curve during rearing for *ad libitum* (AL, —) and food restricted (R, ----) rabbits as described in Rommers *et al.* (2001a).



**Figure 2** Mean daily food intake (g) during rearing for *ad libitum* (AL, —●— no. = 52) and food restricted (R, -○- no. = 52) rabbits. R rabbits were given food to appetite in the last 5 days before insemination. Means  $\pm$  s.d. are presented. An asterisk indicates  $P < 0.05$  for R versus AL rabbits.

exceeded the food intake of AL-rabbits by 34 g/day. Food was provided to the rabbits at 13:00 h.

#### Measurements

**Body weight and food intake.** Animals were weighed individually twice weekly from weaning to first insemination. Food intake was also determined twice weekly by weighing of the feeder at beginning and end of each period and the weight of all food supplies given in between were recorded. For R animals, the amount of food not eaten per day was weighed weekly to determine actual food intake. Based on their actual body weight, food intake was corrected in order to follow the fixed body weight curve. If the daily supply was not eaten completely, remaining food was removed. If wastage occurred, period and cage were recorded, and food intake for this period was scored as a missing value.

**Blood sampling.** At the day of sampling, feeders were closed at 08:00 h. Out of the 52 animals per treatment, 11 animals were randomly taken (11 pairs of full sisters). Post-absorption blood samples were taken weekly from 6 to 17 weeks of age by ear venipuncture between 13:00 and 15:00 h, 5 to 7 h after feeding. Immediately upon collection into tubes containing EDTA, blood samples were centrifuged at 3000 g and plasma stored at  $-20^{\circ}\text{C}$  until assayed for metabolites and hormones.

**Metabolite assays:** glucose, triglycerides (TG), non-esterified fatty acids (NEFA) and plasma urea nitrogen (PUN). Glucose was analysed by the glucose oxidase method using the Glucose Infinity kit from Sigma (Sigma Diagnostic Inc., St Louis, MO, USA). The TG and NEFA concentrations were analysed using enzymatic colorimetric assays from Sigma (Triglycerides Reagent Infinity) and Roche (Roche

Diagnostic GmbH, Mannheim, Germany), respectively. Plasma urea nitrogen was measured by a colorimetric assay using a kit from Dasit (Cornaredo, Italy).

**Hormone assays:** insulin, leptin, total T3, corticosterone. Plasma insulin concentrations were determined by the double antibody/PEG technique using a porcine insulin radio-immuno-assay (RIA) kit (Linco Research Inc., St Charles, MO, USA). The antiserum was guinea pig anti-porcine insulin, while both labelled antigen and standards used purified recombinant human insulin. Leptin concentrations were determined by double antibody RIA using the multi-species leptin kit (Linco Research Inc.). Total T3 was assayed by RIA according to the procedure provided by the manufacturer (Immunotech, Marseille, France). The sensitivity of the assay was 0.13 ng/ml, and the major analogs of T3 did not interfere with the assay. Plasma corticosterone was assayed by RIA, using CORT kit (ICN Biomedicals Inc., Costa Mesa, CA, USA). The sensitivity of the CORT assay was 0.15 ng/ml. Both dilution and recovery tests done on the assay systems for insulin, leptin, T3 and corticosterone, using five different samples of rabbit plasma showed linearity.

#### Statistical analysis

**Body weight, growth rate, food intake, and food efficiency.** Differences between treatments during the subsequent weeks of rearing were tested using the following model:

$$Y_{ij} = \mu + T_i + S_j + (T \times S)_{ij} + e_{ij} \quad (\text{model 1})$$

where  $Y_{ij}$  is dependent variable;  $\mu$  = overall mean;  $T_i$  = treatment R or AL;  $S_j$  = pair of sisters;  $(T \times S)_{ij}$  = interaction between treatment and sister pairs, and  $e_{ij}$  is the residual error. In all analyses, non-significant interactions were deleted from the model. All statistical analyses were performed with the GLM procedure of Statistical Analysis Systems Institute (SAS, 1990). Differences between LSM were analysed by the PDIFF option of the GLM procedure of SAS. Data are presented as means and standard deviations.

**Hormones and metabolites.** Data relative to overall treatment effects during the time-course of the rearing period were analysed by ANOVA (Sokal and Rohlf, 1981) for repeated measurements followed by Student's *t* test. A value of  $P < 0.05$  was considered significant. All statistical analyses were performed using Prism 3.0 software (GraphPad Software, Inc., San Diego, CA, USA).

**Table 1** Body weight, growth rate, food intake, and food efficiency in the restricted and recovery period during rearing for does given food ad libitum (AL, no. = 52) or restricted (R, no. = 52)

	Feeding level			
	<i>Ad libitum</i>		Restricted	
	Mean	s.d.	Mean	s.d.
Restricted period (6 to 12 weeks of age)				
Body weight at start (6 weeks of age) (g)	1295	108	1287	116
Food intake (g/day)	152 <sup>a</sup>	18	106 <sup>b</sup>	6
Growth rate (g/day)	44 <sup>a</sup>	5	29 <sup>b</sup>	2
Food efficiency (g gain/g food)	0.29	0.2	0.27	0.3
Recovery period (12 to 17 weeks of age)				
Body weight at 12 weeks of age (g)	3157 <sup>a</sup>	287	2193 <sup>b</sup>	111
Food intake (g/day)	202 <sup>a</sup>	25	159 <sup>b</sup>	4
Growth rate (g/day)	27 <sup>a</sup>	4	30 <sup>b</sup>	3
Food efficiency (g gain/g food)	0.13	0.9	0.19	0.04
Body weight 17 weeks of age (g)	4101 <sup>a</sup>	376	3563 <sup>b</sup>	135
Body weight at insemination (17.5 weeks) (g)	4202 <sup>a</sup>	388	3798 <sup>b</sup>	220

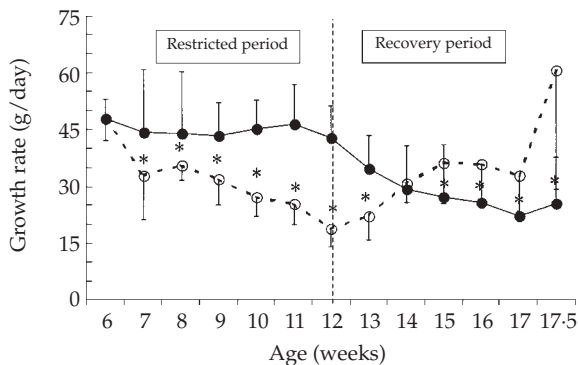
<sup>a,b</sup> Means with a different letter in a row are significantly different ( $P < 0.05$ ).

## Results

### Body weight, growth rate, and food efficiency

Body weight, growth rate, and food efficiency during the restricted and recovery periods are presented in Table 1. Body weight at weaning and at start of the experiment (6 weeks of age) was similar for both treatments. At first insemination (17.5 weeks of age), R rabbits were smaller than AL (3798 (s.d. 220) g and 4202 (s.d. 388) g, respectively;  $P < 0.001$ ).

Growth rate during rearing is presented in Figure 3. Rabbits that were given food *ad libitum* had a high growth rate until approximately 11 weeks of age (between 45 and 48 g/day). Thereafter, body growth decreased and remained constant from 15 weeks of



**Figure 3** Body growth during rearing for *ad libitum* (AL, —●— no. = 52) and food restricted (R, - -○- - no. = 52) rabbits. Means  $\pm$  s.d. are presented. An asterisk indicates  $P < 0.05$  for R versus AL rabbits.

age onwards (between 20 and 25 g/day). Growth rate of the R rabbits decreased to 19 (s.d. 4.9) g/day at 12 weeks of age in the restricted period as a result of the food restriction. During the recovery period, growth rate was aimed to follow the slope of the AL curve (see Figure 1) in order to compensate for the loss in protein (Ledin, 1984) and to stimulate sexual development and puberty (van den Broeck and Lampo, 1977). Increased food intake resulted in an increased growth rate with a growth spurt during the last 5 days before insemination when rabbits were given food to appetite (60 (s.d. 30) g/day).

Food efficiency was affected by feeding level. From weaning to first insemination, *ad libitum* feeding resulted in a lower food efficiency compared with restrictive feeding (0.23 (s.d. 0.23) and 0.26 (s.d. 0.21) for AL and R, respectively;  $P < 0.05$ ).

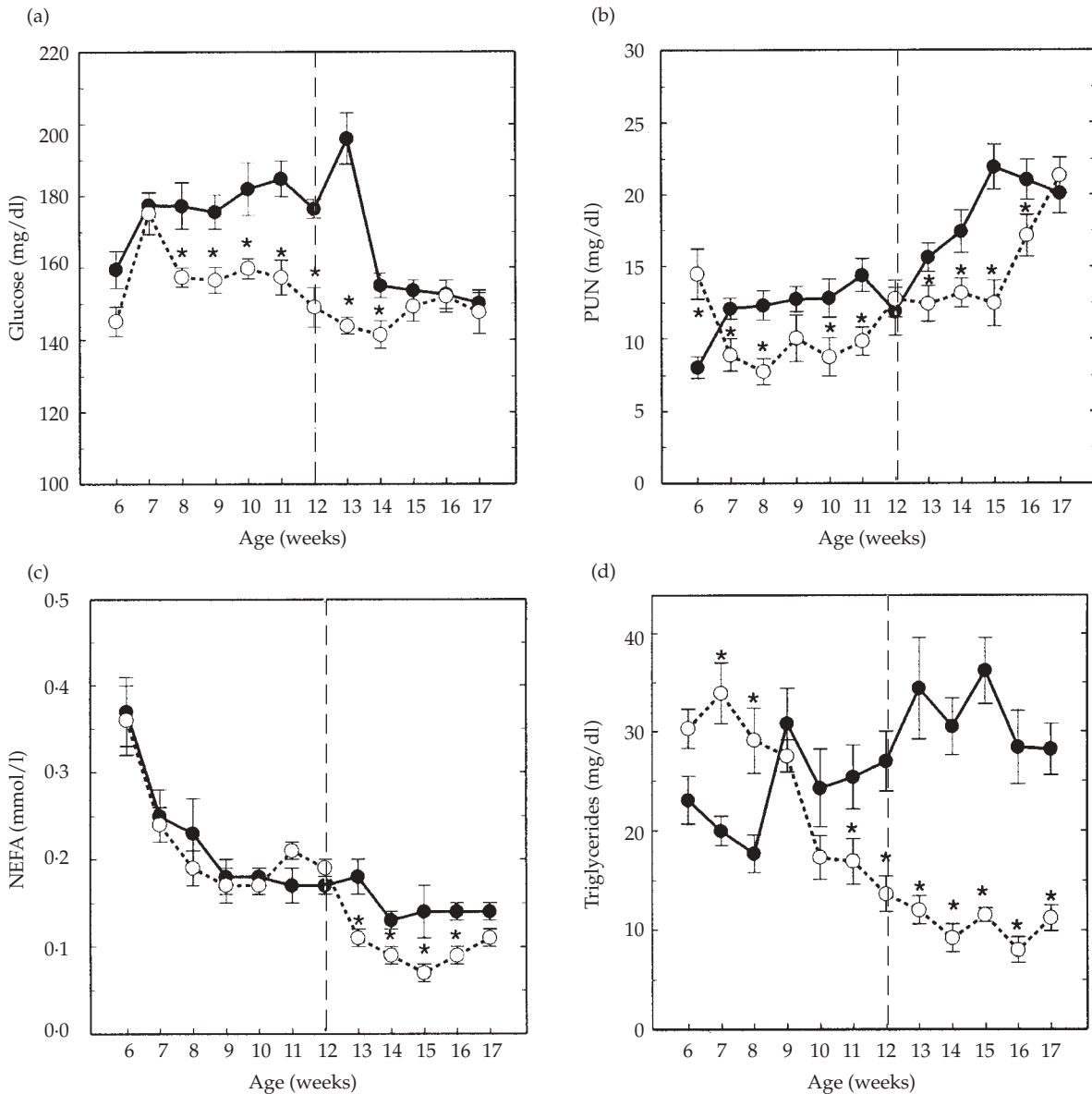
### Metabolic parameters

The plasma profiles of glucose, TG, NEFA, and PUN during the rearing period from week 6 to week 17 of age in AL and R rabbits are shown in Figure 4. Mean plasma glucose concentrations were lower ( $P < 0.05$ ) in R than in AL rabbits from 8 to 14 weeks of age during both the restriction and recovery period (Figure 4a). Only in the last 3 weeks of rearing was glucose restored to control levels in R rabbits, mainly because glucose levels in AL rabbits dropped to R levels.

Triglycerides were higher ( $P < 0.05$ ) in R than in AL rabbits at the beginning of the restriction period from week 7 to week 8 (Figure 4d). From week 11

onwards, TG in R rabbits gradually decreased to values that, at week 17, were almost three-fold lower than those of AL ( $P < 0.05$ ). In AL rabbits, TG increased with advancing age. In both treatments, NEFA showed a similar pattern as their level

dropped ( $P < 0.05$ ) with advancing age, from 0.38 mmol/l at week 6 to about 0.1 mmol/l at week 17 (Figure 4c). In the recovery period from week 13 to 16, NEFA was lower ( $P < 0.05$ ) in R rabbits compared with AL ones. PUN levels were constantly lower

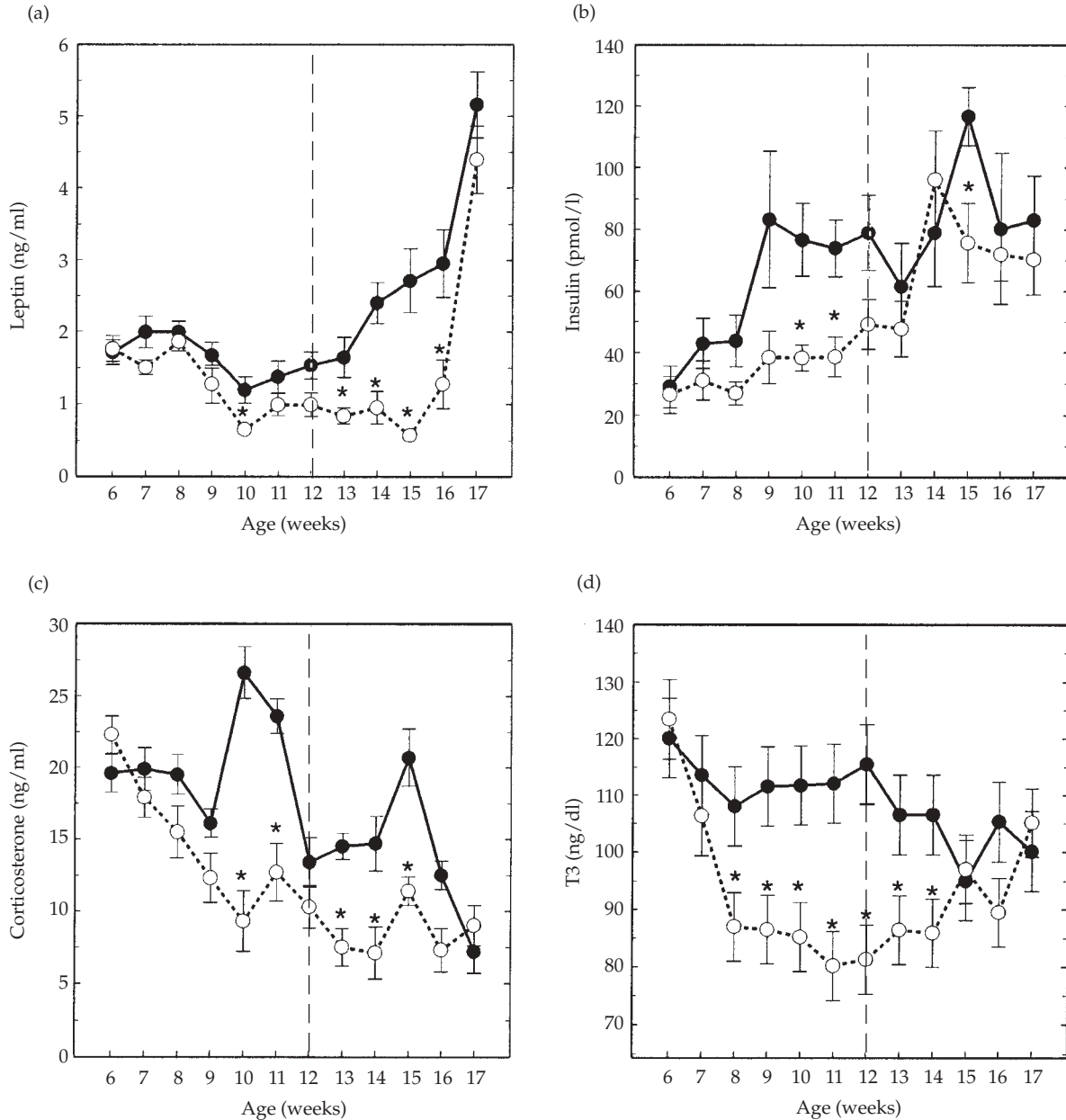


**Figure 4** Metabolic parameters showing (a) glucose, (b) plasma urea nitrogen (PUN), (c) non-esterified fatty acids (NEFA), and (d) triglycerides during the rearing period from week 6 to week 17 of age in rabbits given food *ad libitum* (AL, —●—) or restricted (R, ---○---). The vertical line separates the restriction phase from the following compensatory growth phase. In week 17, R rabbits were given food *ad libitum*. Within each figure, the values, are means  $\pm$  s.e. for 11 animals per treatment group per week. An asterisk indicates  $P < 0.05$  for R versus AL rabbits.

( $P < 0.05$ ) in R compared with AL rabbits, during the restriction as well as in the recovery period (Figure 4b).

#### Hormonal parameters

The blood profiles of immunoreactive leptin, insulin, T3 and corticosterone during the rearing period from



**Figure 5** Endocrine parameters showing immunoreactive (a) leptin, (b) insulin, (c) corticosterone, and (d) tri-iodothyronine (T3) during the rearing period from week 6 to week 17 of age in rabbits given food *ad libitum* (AL, —●—) or restricted (R, .....○.....). The vertical line separates the restriction phase from the following compensatory growth phase. In week 17, R rabbits were given food *ad libitum*. Within each figure, the values, are means  $\pm$  s.e. for 11 animals per treatment group per week. An asterisk indicates  $P < 0.05$  for R versus AL rabbits.

week 6 to week 17 of age in rabbits given food *ad libitum* (AL) or restricted (R) are shown in Figure 5. Leptin levels were similar for both groups until 10 weeks (Figure 5a). From 10 weeks onwards, leptin levels gradually increased in AL, but not in R rabbits. In the last week of rearing, when R rabbits were given food to appetite, leptin concentrations increased five-fold and were similar to that of AL.

An increase in circulating insulin concentrations was observed with advancing age in both AL and R rabbits (Figure 5b). However, insulin was constantly lower in R than AL rabbits during food restriction. In the recovery period, insulin levels of R rabbits rose to values close to those of *ad libitum* animals.

Plasma corticosterone in AL rabbits showed a high variability between successive weeks, but profiles gradually declined with age (Figure 5c). Food restriction reduced circulating corticosterone in R rabbits from 10 until 15 weeks of age. Corticosterone concentrations in the last 2 weeks of rearing were similar for both treatments, due to a sharp decline in AL rabbits.

Compared with AL rabbits, food restriction decreased plasma concentrations of T3 from 8 weeks of age until 14 weeks of age (Figure 5d). During the last 3 weeks of rearing (recovery period) circulating concentrations of T3 were restored to control levels.

## Discussion

In our study, we used an adapted feeding plan in which food intake was reduced from 6 to 12 weeks of age to hinder protein deposition. From 12 weeks of age onwards, rabbits were given increased amounts of food so that they could match growth of rabbits given food *ad libitum*. The purpose of this feeding plan was to obtain a more mature (older aged) rabbit, but prevent the excessive fat deposition that commonly occurs after 12 weeks of age (de Blas *et al.*, 1977; Deltoro and Lopez, 1985). This was done by obtaining the same body weight at 17.5 weeks as that of AL rabbits at 14.5 weeks of age. However, in the recovery period R rabbits were not given food to appetite and metabolic and hormone profiles are not as distinctive as under *ad libitum* feeding conditions. In our study, the general post-absorption state of metabolites and metabolic hormone levels was studied.

The effect of food restriction on the regulatory mechanisms of metabolism during reduced and compensatory growth has been investigated in several animal species (Yambayamba *et al.*, 1996; Rhind *et al.*, 2000), but not in rabbits. Although the basic underlying physiological responses to

underfeeding might be common to all mammals, our data on adjustment of metabolic hormones in the rabbit during long-term food restriction and refeeding are new. Moreover, our energy restriction spanned over a relatively long period of time, from early post weaning up to puberty, during the development of several key organs, such as liver and kidneys (Cantier *et al.*, 1969; Deltoro and Lopez, 1985).

Glucose, the body's major energy source, was lower in R rabbits throughout most of the restriction and the recovery phase due to their lower amount of food intake compared with the AL rabbits. That decreased nutrient intake reduces glucose concentration has also been reported in other species subjected to low planes of nutrition (Booth *et al.*, 1996; Ouellet *et al.*, 2001).

As a consequence of the restricted food intake, plasma immunoreactive insulin was decreased. Interestingly, insulin is the only hormone that in R rabbits was restored to levels found in AL rabbits in the 2nd week of the recovery period. This finding is in agreement with Hornick *et al.* (2000), who stated that normalization of plasma insulin levels was one of the first features occurring within the first days of refeeding, although in our study, plasma insulin levels were restored to control levels later in time.

Insulin is a key player in the control of intermediary metabolism influencing both carbohydrate and lipid metabolism, as well as protein and mineral metabolism. Since there is evidence of active transfer of insulin into the brain, this hormone could have a rôle in signalling the metabolic state of the animal and in the regulation of appetite (Schwartz *et al.*, 1992). Insulin inhibits breakdown of fat in adipose tissue by inhibiting the intracellular lipase that hydrolyses TG to release fatty acids. Insulin facilitates entry of glucose into adipocytes, and within those cells, glucose can be used to synthesize glycerol. This glycerol, along with the fatty acids delivered from the liver, is used to synthesize TG within the adipocyte. By these mechanisms, insulin is involved in further accumulation of TG in fat cells. Low insulin facilitates lypolysis while high insulin stimulates *de novo* lipogenesis (Bruckert and Dejager, 1994; Buczkowska and Jarosz-Chobot, 2001).

Several hormones link growth, metabolism and energy homeostasis (Hornick *et al.*, 2000). Between these, leptin, produced and secreted by adipose cells, is an important component in the long-term regulation of body weight and body fat mass content. In R rabbits, immunoreactive leptin concentrations in plasma were lower compared with

AL rabbits, especially during the compensatory growth phase after week 12. The gradual increase of the circulating leptin with age in AL rabbits may reflect increased adiposity at puberty (around week 12 in rabbit does) probably under the influence of oestrogen as shown in pigs (Barb *et al.*, 2001a). In AL rabbits, an increased fat deposition has been observed from 0.20 to 0.24 of empty body weight between 14.5 and 17.5 weeks of age (Rommers *et al.*, 2001a). Our data on leptin concentrations during food restriction are in agreement with other studies where low nutritional intake prolonged over a longer period induced a similar effect, as demonstrated in pre-pubertal gilts (Barb *et al.*, 2001b), and ruminants (Chilliard *et al.*, 2000).

All other hormone parameters here examined are more related to the body growth or body composition status and are in agreement with other studies in heifers (Yambayamba *et al.*, 1996), bulls (Hornick *et al.*, 1998;) sows (Buonomo and Baile, 1991; Booth *et al.*, 1996; Quesnel *et al.*, 1998) and chickens (Bruggeman *et al.*, 1997).

In the present work, the adrenal axis of R rabbits during both the restriction phase and the subsequent compensatory growth appears to be depressed as corticosterone plasma concentrations were lower compared with AL rabbits. The low corticosterone circulating levels may counteract the excessive catabolic rate of basal metabolism evoked by prolonged food restriction similarly to that found in pre-pubertal gilts by Booth *et al.* (1994). The lower activity of the adrenal axis found in R rabbits during compensatory growth may reflect its responsive adaptation to a long-term stressful condition being applied in the early stage of life (Sapolsky, 1992). Conversely, corticosteroids were found to increase under food restriction in the pig (Prunier and Quesnel, 2000) but decrease in chickens (Zulkifli *et al.*, 1995). These discrepancies may reflect species-specific differences or diverse levels of stress evoked by the bleeding methods as well as dissimilar timing of blood collection, either pre- or post-prandial, in relation to feeding.

In our experimental model, NEFA declined similarly in AL as well as in R rabbits until 12 weeks of age. At the end of the restricted period, food intake of the R rabbits still covered maintenance requirement. Although growth rate was reduced, R rabbits still grew and could not have been in negative energy balance. This implies similar NEFA levels for R and AL rabbits. However, during most of the recovery phase, NEFA were lower in R than in AL rabbits. NEFA are released by the action of hormone sensitive lipase on TG stores in adipose tissue and

increased NEFA concentrations are indicative of negative energy balance (Emery *et al.*, 1992; Rukkwamsuk *et al.*, 1999). Triglycerides were also lower in R rabbits during the recovery period. Hence, decreased circulating NEFA and TG during compensatory growth may reflect redirection of fat metabolism towards reduced lipolysis due to increased plasma insulin concentrations.

The observation that PUN levels were constantly lower in food restricted rabbits throughout the experiment suggests that an alteration of protein metabolism occurred during dietary restriction as well as the recovery period. In energy deficient rabbits, protein-sparing adaptation comes into play to limit turn-over and degradation of protein, whereas, during compensatory growth stimulated by refeeding, increased protein synthesis is associated with decreased nitrogen excretion and lower plasma urea. These findings, which are in agreement with those reported in other species (Sainz *et al.*, 1994; Wykes *et al.*, 1996; Tauson *et al.*, 2002), may provide further evidence for the preferential deposition of lean meat instead of fat tissue in young rabbits undergoing the food restriction (Rommers *et al.*, 2001a).

During food restriction, and for some weeks also during the recovery period, T3 levels were lower than in AL rabbits. Restricted food intake induces animals to spare energy by decreasing basal metabolism (Hornick *et al.*, 2000). Food restriction followed by re-feeding affects thyroidal secretion in cattle (Gerrits *et al.*, 1998; Hornick *et al.*, 1998; Cassar-Makel *et al.*, 2001), swine (Buonomo and Baile, 1991; Booth *et al.*, 1994), and poultry (McMurtry *et al.*, 1988; Buyse *et al.*, 2002). In many species, lower energy intake reduces thyroid function. In our study, T3 concentrations rose to similar levels as AL rabbits after 3 weeks of the recovery period.

In conclusion, long-term nutrient deficiency during development has major neuro-endocrine consequences provoking prominent homeostatic reactions of the corticotropic, leptinergic and thyrotropic axes which persist during compensatory growth, at least up to sexual maturity. During food restriction basal metabolism is reduced owing to the reduction of several anabolic hormones such as T3, and insulin. This pattern of changes in the endocrine status during food restriction is typical and current data in rabbits do confirm those derived from other species, although some specific changes may vary depending on the severity and duration of food restriction. In the recovery period, only insulin adapted within 2 weeks after refeeding. During rearing, hormone and metabolite profiles were



markedly different for the two treatments. Rabbits that were food restricted during rearing as in this study, maintained a higher food intake during first gestation and lactation period and these rabbits produced more kits and milk during the first parity (Rommers *et al.*, 2004). How the metabolic and hormonal profiles during rearing are linked to subsequent voluntary food intake and reproductive performance needs further research.

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